
CROSS-REFERENCE TO RELATED APPLICATIONS

A1
This application is a divisional application of application Serial. No. 09/265,222, filed March 5, 1999, which claims priority under 35 USC § 119(e) to U.S. Provisional Application Serial No. 60/077,618, filed March 10, 1998, the entire contents of which are incorporated herein by reference.

In the claims

Please cancel claims 1-22 and 24-28 without prejudice or disclaimer.

Please amend claim 23 as follows:

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23. (Once Amended) A polynucleotide encoding an apo-B100 protein comprising a proteoglycan receptor⁺ mutation in Site B.

REMARKS

Following entry of this preliminary amendment, claim 23 will be pending in the present application. Claim 23 has been rewritten to eliminate reference to cancelled dependent claims 15-18, and to recite "an apo-B100 protein comprising a proteoglycan receptor⁺ mutation in Site B" (rather than "an apo-B100 protein according to claim 14"). Support for the amendment is found in original claims 14 and 23. No new matter has been added.

Attached hereto is a marked up version of the changes made to the claims and the specification by the current amendment with additions underlined and deletions bracketed. The attached page is captioned "VERSION WITH MARKINGS TO SHOW CHANGES MADE".

In the unlikely event that the Patent Office determines that an extension and/or other relief is required, applicants petition for any required relief including extensions of time and authorizes the Assistant Commissioner to charge the cost of such petitions and/or other fees due in connection with the filing of this document to Deposit Account No. 03-1952 referencing

docket no. 220002059711. However, the Assistant Commissioner is not authorized to charge the cost of the issue fee to the Deposit Account.

Respectfully submitted,

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VERSION WITH MARKINGS TO SHOW CHANGES MADE

U.S. Serial No. To Be Assigned
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In the specification

On page 1, under the title, a new paragraph has been added.

In the claims

23. (Once Amended) A polynucleotide encoding an apo-B100 protein [according to any one of claims 14 to 18] comprising a proteoglycan receptor⁺ mutation in Site B.